

Food and Drug Administration 10903 New Hampshire Avenue Document Control Center – WO66-G609 Silver Spring, MD 20993-0002

February 25, 2016

GE Healthcare % Ms. Tracey Ortiz Regulatory Affairs Director 9900 W. Innovation Drive WAUWATOSA WI 53226

Re: K160184

Trade/Device Name: Voluson S6, Voluson S8, Voluson S10

Regulation Number: 21 CFR 892.1550

Regulation Name: Ultrasonic pulsed doppler imaging system

Regulatory Class: II

Product Code: IYN, IYO, ITX Dated: January 20, 2016 Received: January 27, 2016

Dear Ms. Ortiz:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638 2041 or (301) 796-7100 or at its Internet address

http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to

<u>http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm</u> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm.

Sincerely yours,

Robert Ods

Robert Ochs, Ph.D.
Director
Division of Radiological Health
Office of In Vitro Diagnostics
and Radiological Health
Center for Devices and Radiological Health

Enclosure



510(k) Premarket Notification Submission

DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration	Form Approved: OMB No. 0910-0120 Expiration Date: January 31, 2017
Indications for Use	See PRA Statement on last page.
510(k) Number (if known)	
K160184	
Device Name Voluson S6 / Voluson S8 / Voluson S10	
Indications for Use (Describe)	
The device is a general-purpose ultrasound system. Specific clinical applications and exam Abdominal (including renal and GYN/pelvic); Pediatric; Small Organ (breast, testes, thyroid and neonatal patients); Neonatal Cephalic; Adult Cephalic; Cardiac (adult and pediatric); Pediatric); Pediatric (TR); Transvaginal (TV).	l, salivary gland, lymph nodes, pediatric
Type of Use (Select one or both, as applicable)	
, , , , , , , , , , , , , , , , , , ,	nter Use (21 CFR 801 Subpart C)
	(
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FOR FDA USE ONLY	
Concurrence of Center for Devices and Radiological Health (CDRH) (Signature)	

FORM FDA 3881 (1/14) Page 1 of 2 PSC Publishing Services (301) 443-6740 EF



510(k) Premarket Notification Submission

This section applies only to requirements of the Paperwork Reduction Act of 1995.

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Department of Health and Human Services Food and Drug Administration Office of Chief Information Officer Paperwork Reduction Act (PRA) Staff PRAStaff@fda.hhs.gov

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510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 Ultrasound System

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Me	ode of Ope	eration				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler	Power Doppler	Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes]
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P	P	P	P	P	P	P	P	[5,6,9]
Abdominal ^[1]	P	P	P	P	P	P	P	P	P	P	[5,6,9]
Pediatric	P	P	P	P	P	P	P	P	P	P	[5,6,9]
Small Organ ^[2]	P	P	P	P	P	P	P	P	P	P	[5,6,9]
Neonatal Cephalic	P	P	P	P	P	P	P	P	P	P	[5]
Adult Cephalic	P	P	P	P	P	P	P	P	P	P	
Cardiac ^[3]	P	P	P	P	P	P	P	P	P	P	[5]
Peripheral Vascular	P	P	P		P	P	P	P	P	P	[5,6,9]
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[5,6,9]
Musculo-skeletal Superficial	P	P	P		P	P	P	P	P	P	[5,6,9]
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]	P	P	P		P	P	P	P	P	P	[5,6,9]
Transvaginal	P	P	P		P	P	P	P	P	P	[5,6,9]
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric
- [5] 3D/4D Imaging Mode
- [6] Includes imaging of guidance of biopsy (2D/3D/4D)
- [7] Includes infertility monitoring of follicle development
- [8] Includes urology/prostate
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with IC9-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mo	de of Oper	ation				
Clinical Application	В	M	PW Doppler	CW Doppler	Color Doppler	Color M Doppler		Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Anatomy/Region of Interest Ophthalmic			Боррісі	Боррісі	Боррісі	Doppier	Doppiei	Wiodes	maging	1 tilse	[Ivotes)
*											
Fetal / Obstetrics ^[7]	N	N	N		N	N	N	N	N	N	[6,9]
Abdominal ^[1]	N	N	N		N	N	N	N	N	N	[6,9]
Pediatric											
Small Organ ^[2]											
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular											
Musculo-skeletal Conventional											
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]	N	N	N		N	N	N	N	N	N	[6,9]
Transvaginal ^[10]	N	N	N		N	N	N	N	N	N	[6,9]
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric
- [5] 3D/4D Imaging Mode
- [6] Includes imaging of guidance of biopsy (2D/3D/4D)
- [7] Includes infertility monitoring of follicle development
- [8] Includes urology/prostate
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with 4C-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mod	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	M	PW Doppler	CW Doppler	Color Doppler	Color M Doppler		Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P		P	P	P	P	P	P	[6]
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	[6]
Pediatric	P	P	P		P	P	P	P	P	P	[6]
Small Organ ^[2]											
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular	P	P	P		P	P	P	P	P	P	[6]
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[6]
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with C1-5-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mod	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler		Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P		P	P	P	P	P	P	[6]
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	[6]
Pediatric	P	P	P		P	P	P	P	P	P	[6]
Small Organ ^[2]											
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular											
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[6]
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with 8C-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mo	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler	Power Doppler	Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P		P	P	P	P	P	P	
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	
Pediatric	P	P	P		P	P	P	P	P	P	
Small Organ ^[2]											
	P	P	P		P	P	P	P	P	P	
Neonatal Cephalic	P	P	P		P	P	P	P	P	P	
Adult Cephalic											
Cardiac ^[3]	P	P	P		P	P	P	P	P	P	
Peripheral Vascular	P	P	P		P	P	P	P	P	P	
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	
Musculo-skeletal Superficial	P	P	P		P	P	P	P	P	P	
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic N = powindication: P = provious											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with E8C-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

			_		Mo	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler	Power Doppler		Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P		P	P	P	P	P	P	[6,9]
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	[6,9]
Pediatric											
Small Organ ^[2]											
Neonatal Cephalic	P	P	P		P	P	P	P	P	P	[6,9]
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular											
Musculo-skeletal Conventional											
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]	P	P	P		P	P	P	P	P	P	[6,9]
Transvaginal	P	P	P		P	P	P	P	P	P	[6,9]
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric
- [5] 3D/4D Imaging Mode
- [6] Includes imaging of guidance of biopsy (2D/3D/4D)
- [7] Includes infertility monitoring of follicle development
- [8] Includes urology/prostate
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with with 12L-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mod	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler	Power Doppler	Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]											
Abdominal ^[1]											
Pediatric	P	P	P		P	P	P	P	P	P	[6,9]
Small Organ ^[2]	P	P	P		P	P	P	P	P	P	[6,9]
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular	P	P	P		P	P	P	P	P	P	[6,9]
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[6,9]
Musculo-skeletal Superficial	P	P	P		P	P	P	P	P	P	[6,9]
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with 9L-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mo	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler	Power Doppler	Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P		P	P	P	P	P	P	[6]
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	[6]
Pediatric	P	P	P		P	P	P	P	P	P	[6]
Small Organ ^[2]	P	P	P		P	P	P	P	P	P	[6]
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular	P	P	P		P	P	P	P	P	P	[6]
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[6]
Musculo-skeletal Superficial	P	P	P		P	P	P	P	P	P	[6]
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with 3Sc-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mod	de of Oper	ration				
Clinical Application Anatomy/Region of Interest	В	M	PW Doppler	CW Doppler	Color	Color M Doppler		Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P	P	P	P	P	<u>P</u>	P	P	[6]
Abdominal ^[1]	P	P	P	P	P	P	P	P	P	P	[6]
Pediatric	P	P	P	P	P	P	P	P	P	P	[6]
Small Organ ^[2]											
Neonatal Cephalic											
Adult Cephalic	P	P	P	P	P	P	P	P	P	P	[6]
Cardiac ^[3]	P	P	P	P	P	P	P	P	P	P	[6]
Peripheral Vascular											
Musculo-skeletal Conventional											
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



510(k) Premarket Notification Submission

Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with RIC5-9A-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mod	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color Doppler	Color M Doppler	Power Doppler	Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	N	N	N		N	N	N	N	N	N	[5,6,9]
Abdominal ^[1]	N	N	N		N	N	N	N	N	N	[5,6,9]
Pediatric											
Small Organ ^[2]											
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular											
Musculo-skeletal Conventional											
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]	N	N	N		N	N	N	N	N	N	[5,6,9]
Transvaginal	N	N	N		N	N	N	N	N	N	[5,6,9]
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Diagnostic Ultrasound Indications for Use Form

Voluson S6 / Voluson S8 / Voluson S10 with RAB6-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

					Mod	de of Oper	ation				
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color	Color M Doppler		Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]	P	P	P		P	P	P	P	P	P	[5,6]
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	[5,6]
Pediatric	P	P	P		P	P	P	P	P	P	[5,6]
Small Organ ^[2]											
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular											
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[5,6]
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Diagnostic Ultrasound Indications for Use Form

Voluson S8 / Voluson S10 with ML6-15-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

	Mode of Operation										
Clinical Application	В	M	PW	CW	Color	Color M	Power	Combined		Coded	Other
Anatomy/Region of Interest			Doppler	Doppler	Doppler	Doppler	Doppler	Modes*	Imaging	Pulse	[Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]											
Abdominal ^[1]	P	P	P		P	P	P	P	P	P	[6,9]
Pediatric	P	P	P		P	P	P	P	P	P	[6,9]
Small Organ ^[2]	P	P	P		P	P	P	P	P	P	[6,9]
Neonatal Cephalic											
Adult Cephalic											
Cardiac ^[3]											
Peripheral Vascular	P	P	P		P	P	P	P	P	P	[6,9]
Musculo-skeletal Conventional	P	P	P		P	P	P	P	P	P	[6,9]
Musculo-skeletal Superficial	P	P	P		P	P	P	P	P	P	[6,9]
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^{[8}											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Diagnostic Ultrasound Indications for Use Form

Voluson S8 / Voluson S10 with 12S-RS Transducer

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

	Mode of Operation										
Clinical Application Anatomy/Region of Interest	В	М	PW Doppler	CW Doppler	Color	Color M Doppler		Combined Modes*	Harmonic Imaging	Coded Pulse	Other [Notes)
Ophthalmic											
Fetal / Obstetrics ^[7]											
Abdominal ^[1]											
Pediatric	P	P	P	P	P	P	P	P	P	P	
Small Organ ^[2]	P	P	P	P	P	P	P	P	P	P	
Neonatal Cephalic	P	P	P	P	P	P	P	P	P	P	
Adult Cephalic	P	P	P	P	P	P	P	P	P	P	
Cardiac ^[3]	P	P	P	P	P	P	P	P	P	P	
Peripheral Vascular											
Musculo-skeletal Conventional											
Musculo-skeletal Superficial											
Other											
Exam Type, Means of Access											
Transesophageal											
Transrectal ^[8]											
Transvaginal											
Transuretheral											
Intraoperative											
Intraoperative Neurological											
Intravascular											
Laparoscopic											

N = new indication; P = previously cleared by FDA

Notes: [1] Abdominal includes renal, GYN/Pelvic

- [2] Small organ includes breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients
- [3] Cardiac is Adult and Pediatric.
- [5] 3D/4D Imaging Mode.
- [6] Includes imaging of guidance of biopsy (2D/3D/4D).
- [7] Includes infertility monitoring of follicle development.
- [8] Includes urology/prostate.
- [9] Elastography imaging- Elasticity
- [*] Combined modes are B/M, B/Color M, B/PWD, B/Color/PWD, B/PWD

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Concurrence of CDRH, Office of In Vitro Diagnostics and Radiological Health (OIR)



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510(k) Summary

In accordance with 21 CFR 807.92 the following summary of information is provided:

<u>Date:</u> January 20, 2016

Submitter: GE Healthcare

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Device: Trade Name: Voluson S6 / Voluson S8 / Voluson S10

Common/Usual Name: Ultrasound system

<u>Classification Names:</u> Class II

Product Code: Ultrasonic Pulsed Doppler Imaging System. 21CFR 892.1550 90-IYN

Ultrasonic Pulsed Echo Imaging System, 21CFR 892.1560, 90-IYO

Diagnostic Ultrasound Transducer, 21 CFR 892.1570, 90-ITX

Predicate Device(s): K141639 Voluson S Series; Voluson S6/S8 Diagnostic

Ultrasound System

K152567 Voluson E Series; E6/E8/E10 Diagnostic Ultrasound

System

K142472 Voluson E Series; E6/E8/E10 Diagnostic Ultrasound

System

<u>Device Description:</u> The systems are full-featured Track 3 ultrasound systems,

primarily for general radiology use and specialized for OB/GYN with particular features for realtime 3D/4D acquisition. They consist of a mobile console with keyboard control panel; color LCD/TFT touch panel, color video display and optional image storage and printing devices. They provide high performance ultrasound imaging and analysis and have comprehensive networking and DICOM capability. They utilize a variety of linear, curved linear, matrix phased array transducers including



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mechanical and electronic scanning transducers, which provide accurate realtime three dimensional imaging supporting all standard acquisition modes.

Intended Use:

The device is a general purpose ultrasound system. Specific clinical applications remain the same as previously cleared: Fetal/OB; Abdominal (including renal and GYN/pelvic); Pediatric; Small Organ (breast, testes, thyroid, salivary gland, lymph nodes, pediatric and neonatal patients); Neonatal and Adult Cephalic; Cardiac (adult and pediatric); Peripheral Vascular (PV); Musculo-skeletal Conventional and Superficial; Transrectal (TR); Transvaginal (TV).

Technology:

The Voluson S6 / Voluson S8 / Voluson S10 employs the same fundamental scientific technology as its predicate devices.

<u>Determination of</u> <u>Substantial Equivalence:</u>

Comparison to Predicates

The Voluson S6 / Voluson S8 / Voluson S10 is substantially equivalent to the predicate devices with regards to intended use, imaging capabilities, technologicial characteristics and safety and effectiveness.

- The systems are all intended for diagnostic ultrasound imaging and fluid flow analysis.
- The Voluson S6/S8/S10 and predicate Voluson S6/S8 systems have the same clinical intended use.
- The Voluson S6/S8/S10 and predicate Voluson S6/S8 systems have the same imaging modes.
- The Voluson S6/S8/S10 and predicate Voluson S6/S8 systems transducers are identical except for the removal of six transducers (RAB4-8-RS, RIC5-9W-RS, P2D, RAB2-5-RS, AB2-7-RS, RAB2-6-RS); the removal of ML6-15-RS and the 12S-RS probes on Voluson S6; the addition of two new transducers IC9-RS (similar to IC9-D on predicate Voluson E Series K152567), and RIC5-9A-RS (similar to RIC5-9W-RS on predicate Voluson S Series K141639)
- The Voluson S6/S8/S10 and predicate Voluson S6/S8 systems have the same indications for use.
- The systems are manufactured with materials which have been evaluated and found to be safe for the intended use



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of the device.

- The systems have acoustic power levels which are below the applicable FDA limits.
- The Voluson S6/S8/S10 and predicate Voluson S6/S8 systems have similar capability in terms of performing measurements, capturing digital images, reviewing and reporting studies.
- The Voluson S6/S8/S10 and predicate systems have been designed in compliance with approved electrical and physical safety standards.
- The Voluson S6/S8/S10 adds a SIM card to the system to allow users to send images and text via MMS_SMS wirelessly. (previously cleared with K142472).
- The Voluson S6/S8/S10 adds a GYN measurement tool created by IOTA Group called IOTA LR2 model (previously cleared with K142472).
- The proposed Voluson S6/S8/S10 adds software features called SonoAVC antral and SonoAVC general. (previously cleared with K152567).
- The proposed Voluson S6/S8/S10 beam former has changed to STB MLA4 previously cleared with K152567 / K142472.

Summary of Non-Clinical Tests:

The device has been evaluated for acoustic output, biocompatibility, cleaning and disinfection effectiveness as well as thermal, electrical, electromagnetic, and mechanical safety, and has been found to conform to applicable medical device safety standards. The Voluson S6/S8/S10 systems and its applications comply with voluntary standards:

- AAMI/ANSI ES60601-1, Medical Electrical Equipment Part 1: General Requirements for Safety
- IEC60601-1-2, Medical Electrical Equipment Part 1-2:General Requirements for Safety – Collateral Standard: Electromagnetic Compatibility Requirements and Tests
- IEC60601-2-37, Medical Electrical Equipment Part 2-37:Particular Requirements for the Safety of Ultrasonic Medical Diagnostic and Monitoring Equipment
- NEMA UD 3, Standard for Real Time Display of Thermal



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and Mechanical Acoustic Output Indices on Diagnostic Ultrasound Equipment

- ISO10993-1, Biological Evaluation of Medical Devices-Part 1: Evaluation and Testing
- NEMA UD 2, Acoustic Output Measurement Standard for Diagnostic Ultrasound Equipment
- ISO14971, Application of risk management to medical devices
- NEMA, Digital Imaging and Communications in Medicine (DICOM) Set. (Radiology)

The following quality assurance measures were applied to the development of the system:

- Risk Analysis
- Requirements Reviews
- Design Reviews
- Testing on unit level (Module verification)
- Integration testing (System verification)
- Performance testing (Verification)
- Safety testing (Verification)

Transducer materials and other patient contact materials are biocompatible.

Summary of Clinical Tests:

The subject of this premarket submission, Voluson S6 / Voluson S8 / Voluson S10, did not require clinical studies to support substantial equivalence.

Conclusion: GE Healthcare considers the Voluson S6 / Voluson S8 / Voluson S10 to be as safe, as effective, and performance is substantially equivalent to the predicate device(s).